

APPLICATION NUMBER	FILED RECEIPT DATE	FIRST NAMED APPLICANT	ATTORNEY DOCKET NUMBER
10/085,378	02/27/2002	Sean D. Monahan	.013.04.05

[illegible]

Date Mailed: 10/30/2002

Filing Date Granted

The required item(s) identified below must be timely submitted to avoid abandonment:


- A substitute specification in compliance with 37 CFR 1.52, 1.121(b)(3), and 1.125, is required. The specification, claims, or abstract page(s) submitted is not acceptable and cannot be scanned or properly stored because:
 - The line spacing on the specification, claims, or abstract is not 1-1/2 or double spaced (see 37 CFR 1.52(b)).
- This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to the final rulemaking notice published at 55 FR 18230 (May 1, 1990), and 1114 OG 29 (May 15, 1990). If the effective filing date is on or after July 1, 1998, see the final rulemaking notice published at 63 FR 29620 (June 1, 1998) and 1211 OG 82 (June 23, 1998). If the effective filing date is on or after September 8, 2000, see the final rulemaking notice published in the Federal Register at 65 FR 54604 (September 8, 2000) and 1238 OG 145 (September 19, 2000). Applicant must provide an initial computer readable form (CRF) copy of the "Sequence Listing", an initial paper or compact disc copy of the "Sequence Listing", as well as an amendment directing its entry into the application. Applicant must also provide a statement that the content of the sequence listing information recorded in computer readable form is identical to the written (on paper or compact disc) sequence listing and, where applicable, includes no new matter as required by 37 CFR 1.821, 1.821(f), 1.821(g), 1.825, and 1.825(f).

...the fact that the *in vitro* and *in vivo* results are in good agreement.

For questions regarding compliance to these requirements, please contact:

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A copy of this notice MUST be returned with the reply.



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